

EXHIBIT B

(Redacted as Described in the Motion to Seal)

1 IN THE UNITED STATES DISTRICT COURT
2 FOR THE MIDDLE DISTRICT OF NORTH CAROLINA

3
4 SYNGENTA CROP PROTECTION, LLC

5 Plaintiff

6 vs.

CIVIL ACTION NO:

7 WILLOWOOD, LLC, et al.

1:15-CV-274

8 Defendants

9 _____/

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13 The Videodeposition of JOSEPH M.D.
14 FORTUNAK, Ph.D. was held on Monday, October 10, 2016,
15 commencing at 9:00 a.m., at Kirkland & Ellis, LLP, 655
16 15th Street, N.W., Suite 1200, Washington D.C. 20005,
17 before Christina Essi Pagano, Notary Public.

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20
21 Reported By: Christina Essi Pagano, RPR

<p>ATTORNEYS' EYES ONLY</p> <p>Page 78</p> <p>1</p> <p>2</p>	<p>ATTORNEYS' EYES ONLY</p> <p>Page 80</p> <p>1 during the condensation step of manufacture of</p> <p>2 azoxystrobin supplied by TaiHe to Willowood, correct?</p> <p>3 A. That's correct.</p> <p>4 Q.</p>
<p>ATTORNEYS' EYES ONLY</p> <p>Page 79</p> <p>1</p> <p>4 TaiHe documentation, I think it would be very, very</p> <p>5 difficult to reproduce this in order to see what levels</p> <p>6 would regularly project into an isolated azoxystrobin</p> <p>7 technical.</p> <p>8 Q. And '761 patent requires that the</p> <p>9 condensation step -- and can we just agree that we'll</p> <p>10 call it the condensation step for purposes of this</p> <p>11 discussion?</p> <p>12 A. Yes.</p> <p>13 Q. The '761 patent requires that the</p> <p>14 condensation step be done in the presence of between .1</p> <p>15 molar percent and 2.0 molar percent DABCO, correct?</p> <p>16 A. That's right.</p> <p>17 Q. And from this testing performed by CAC</p> <p>18 Shanghai, you could not quantify the amount of</p> <p>19 azoxystrobin -- strike that.</p> <p>20 From this testing done by CAC Shanghai, you</p> <p>21 could not quantify the amount of DABCO that was present</p>	

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12 Q. But the CAC testing you weren't able to
13 quantify how much DABCO was used in the manufacturing
14 process, correct?
15 A. That's correct.
16 Q. And from the Syngenta testing you were not
17 able to quantify how much DABCO was used in the
18 manufacturing process, correct?
19 A. That's correct.
20 Q.

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8 Q. But commercially reasonable companies do
9 engage in design-around efforts, to the best of your
10 knowledge, correct?
11 MS. BALTZER: Objection, foundation.
12 THE WITNESS: That's correct.
13

ATTORNEYS' EYES ONLY Page 118 1	ATTORNEYS' EYES ONLY Page 120 1 sold to Willowood? 2 MS. BALTZER: Objection, foundation. 3 THE WITNESS: In terms of how much it 4 contributes to the cost of the raw materials, it won't 5 be a large contribution. 6
ATTORNEYS' EYES ONLY Page 119 1 6 Q. So after the quotation from the '761 patent 7 that you cite to, you say, quote, given the information 8 in the '761 patent, it would not be commercially 9 reasonable for Willowood's manufacturer, (TaiHe) to use 10 more DABCO than 2 molar percent, which would add to the 11 manufacturing expense without any significant 12 improvement in yield. 13 Do you see that? 14 A. Yes. 15 Q. What is the expense of DABCO? What does it 16 cost? 17 A. Not terrible expensive. 18 Q. Okay. So increasing DABCO from, for 19 example, 1.9 molar percent to 2.1 molar percent, do you 20 know how much that would increase the cost of 21 manufacturing the azoxystrobin produced by TaiHe and	

ATTORNEYS' EYES ONLY Page 126 1	ATTORNEYS' EYES ONLY Page 128 1
ATTORNEYS' EYES ONLY Page 127 1 5 BY MR. TILLER: 6 Q. Okay. When you used the term increase 7 toxicity, to whom would it be toxic? 8 A. It could be toxic either to the plants that 9 to is used to be applied to or it could be toxic to 10 people who use the agriculture end product that comes 11 from the plant. 12 Q. What are the toxic effects? 13 A. I don't recall. 14 Q. Putting aside the amount of DABCO used in 15 the manufacturing process, how much of this residue 16 would have to be in the azoxystrobin product that was 17 applied to the plants, how much residue would have to 18 be in that product in order for it to be toxic? 19 MS. BALTZER: Objection, foundation. 20 THE WITNESS: I do not know that that has 21 been determined.	ATTORNEYS' EYES ONLY Page 129 1

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8 BY MR. TILLER:

9 Q. So when you say 723, we're looking at
10 Weintritt?

11 A. Yeah, at Weintritt.

12 Q. Okay.

13 A. And if I compare Weintritt with a compound
14 patent for azoxystrobin, which I believe is the '076,
15 both of those cover millions and millions of compounds.
16 And the structure azoxystrobin would be covered under
17 both of those.

18 Q. Under '076 and Weintritt?

19 A. Yeah.

20 Q.

ATTORNEYS' EYES ONLY Page 162 1	ATTORNEYS' EYES ONLY Page 164 1 12 Q. Well, Weintritt is claiming the synthesis 13 of azoxystrobin in the presence of 2 to 40 molar 14 percent of DABCO, correct? 15 MS. BALTZER: Objection, foundation. 16 THE WITNESS: Looking at the scope of 17 Weintritt's claims, one could infer that -- one could 18 say that this is contemplated. It's not exemplified, 19 nor is it explicitly laid out. It said, no, I can make 20 azoxystrobin with 2 mole percent of DABCO. 21 BY MR. TILLER:
ATTORNEYS' EYES ONLY Page 163 1 Q. Okay. So I thought we had agreed that 2 Weintritt claimed a process for preparing compounds of 3 the general formula one and we agreed that general 4 formula one includes, among many other compounds, but 5 it includes azoxystrobin, correct? 6 A. That's correct. 7 MS. BALTZER: Objection, foundation. 8 BY MR. TILLER: 9 Q. That's correct, correct? 10	ATTORNEYS' EYES ONLY Page 165 1 Q. So you don't think the fact that Weintritt 2 claims the production or the manufacture of 3 azoxystrobin in the presence of 2 to 40 molar percent, 4 you don't think that despite that claim, that a person 5 of ordinary skill in the art would not read this to 6 disclose the manufacture of azoxystrobin in the 7 presence of 2 to 40 molar percent of DABCO? 8 MS. BALTZER: Objection, foundation, asked 9 and answered. 10 THE WITNESS: I'm going to answer you as a 11 chemist. You said the word manufacturing. 12 Manufacturing means that I'm making something and I can 13 make it, I can make it efficiently, and I can sell it. 14 The scope of Weintritt's claims would cover the 15 manufacture of 2 mole -- of azoxystrobin using 16 different components that he hasn't exemplified the use 17 of in his examples, and using down to 2 mole percent of 18 DABCO. So within that -- so using 2 mole percent of 19 DABCO and using a condensation to make azoxystrobin in 20 that way would fall under the claims of Weintritt. 21 BY MR. TILLER:

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12 Q. Okay. So still, sitting here today, you
13 cannot conclusively say how much DABCO was used during
14 the manufacture of the azoxystrobin technical included
15 in that sample of Azoxy 2SC, correct?
16 MS. BALTZER: Objection, form.
17 THE WITNESS: So the best available
18 information you can get is to measure the presence of
19 DABCO, the byproducts derived from it, and that doesn't
20 allow you to quantify how much DABCO was used.
21 BY MR. TILLER:

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1 Q. Understood. Actually, the best available
2 information would be go and take samples of the
3 azoxystrobin technical during the manufacturing and
4 then test that, correct?
5 MS. BALTZER: Objection, form.
6 THE WITNESS: Correct, but that hasn't been
7 offered as an option by Willowood or TaiHe.
8 BY MR. TILLER:
9 Q. Hasn't been asked, hasn't been asked, has
10 it?
11 MS. BALTZER: Objection, form.
12 THE WITNESS: I don't know.
13 BY MR. TILLER:
14 Q.